

STN Columbus

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NEWS 5 DEC 14 2006 MeSH terms loaded for MEDLINE file segment of TOXCENTER
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NEWS 7 DEC 21 IPC search and display fields enhanced in CA/Caplus with the
IPC reform
NEWS 8 DEC 23 New IPC8 SEARCH, DISPLAY, and SELECT fields in USPATFULL/
USPAT2
NEWS 9 JAN 13 IPC 8 searching in IFIPAT, IFIUDb, and IFICDB
NEWS 10 JAN 13 New IPC 8 SEARCH, DISPLAY, and SELECT enhancements added to
INPADOC
NEWS 11 JAN 17 Pre-1988 INPI data added to MARPAT
NEWS 12 JAN 17 IPC 8 in the WPI family of databases including WPIFV
NEWS 13 JAN 30 Saved answer limit increased
NEWS 14 JAN 31 Monthly current-awareness alert (SDI) frequency
added to TULSA

NEWS EXPRESS FEBRUARY 15 CURRENT VERSION FOR WINDOWS IS V8.01a,
CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
AND CURRENT DISCOVER FILE IS DATED 19 DECEMBER 2005.
V8.0 AND V8.01 USERS CAN OBTAIN THE UPGRADE TO V8.01a AT
<http://download.cas.org/express/v8.0-Discover/>

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* * * * * STN Columbus * * * * *

FILE 'HOME' ENTERED AT 11:32:18 ON 18 FEB 2006

=> FILE REG

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	3.57	3.57

FILE 'REGISTRY' ENTERED AT 11:42:45 ON 18 FEB 2006

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STRUCTURE FILE UPDATES: 16 FEB 2006 HIGHEST RN 874442-49-6
DICTIONARY FILE UPDATES: 16 FEB 2006 HIGHEST RN 874442-49-6

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH January 6, 2006

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*****
*
* The CA roles and document type information have been removed from *
* the IDE default display format and the ED field has been added,   *
* effective March 20, 2005. A new display format, IDERL, is now    *
* available and contains the CA role and document type information. *
*
*****
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Structure search iteration limits have been increased. See HELP SLIMITS
for details.

REGISTRY includes numerically searchable data for experimental and
predicted properties as well as tags indicating availability of
experimental property data in the original document. For information
on property searching in REGISTRY, refer to:

<http://www.cas.org/ONLINE/UG/regprops.html>

=>

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L1 STRUCTURE UPLOADED

=> s l1

SAMPLE SEARCH INITIATED 11:42:55 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 3714 TO ITERATE

53.9% PROCESSED 2000 ITERATIONS 1 ANSWERS
INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**

PROJECTED ITERATIONS: 70626 TO 77934

PROJECTED ANSWERS: 1 TO 118

L2 1 SEA SSS SAM L1

=> s l1 sss full

FULL SEARCH INITIATED 11:43:07 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 75735 TO ITERATE

100.0% PROCESSED 75735 ITERATIONS 198 ANSWERS
SEARCH TIME: 00.00.02

L3 198 SEA SSS FUL L1

=> file caplus

COST IN U.S. DOLLARS

SINCE FILE
ENTRY

TOTAL
SESSION

STN Columbus

FULL ESTIMATED COST

166.94

170.51

FILE 'CAPLUS' ENTERED AT 11:43:17 ON 18 FEB 2006

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FILE COVERS 1907 - 18 Feb 2006 VOL 144 ISS 9

FILE LAST UPDATED: 17 Feb 2006 (20060217/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

<http://www.cas.org/infopolicy.html>

=> s l3

L4 7 L3

=> d l4 1-7 bib abs fhitr

L4 ANSWER 1 OF 7 CAPLUS COPYRIGHT 2006 ACS on STN

Full Text

AN 2005:980814 CAPLUS

DN 143:431985

TI Privileged structure-based ligands for melanocortin receptors-tetrahydroquinolines, indoles, and aminotetralines

AU Fisher, Matthew J.; Backer, Ryan T.; Husain, Saba; Hsiung, Hansen M.; Mullaney, Jeffrey T.; O'Brian, Thomas P.; Ornstein, Paul L.; Rothhaar, Roger R.; Zgombick, John M.; Briner, Karin

CS Lilly Research Laboratories, A Division of Eli Lilly and Company, Lilly Corporate Center, Indianapolis, IN, 46258, USA

SO Bioorganic Medicinal Chemistry Letters (2005), 15(20), 4459-4462

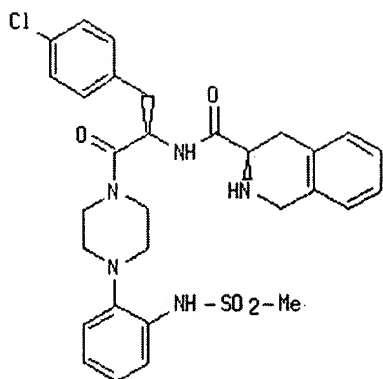
CODEN: BMCLE8; ISSN: 0960-894X

PB Elsevier B.V.

DT Journal

LA English

GI



AB Substitution of the aryl sulfonamide moiety contained in MC4 agonist (I) with bicyclic heterocycles and aminotetralines produced compds. with MC4 activity. The heterocycles represent alternative privileged structures to that contained in I. Compds. in which the polar group of the privileged structure was displayed in an endocyclic fashion were not as active as the parent agonist I, while those with an exocyclic polar group afforded activity competitive with I.

IT 444619-63-0P

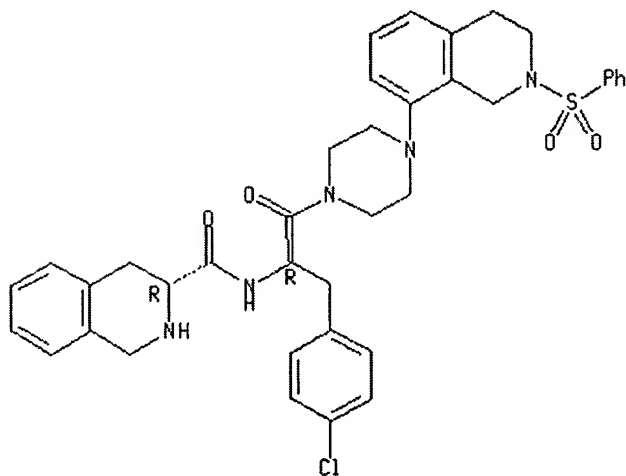
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(privileged structure-based ligands for melanocortin receptors-tetrahydroquinolines, indoles, and aminotetralines)

RN 444619-63-0 CAPLUS

CN 3-Isoquinolinecarboxamide, N-[(1R)-1-[(4-chlorophenyl)methyl]-2-oxo-2-[4-[1,2,3,4-tetrahydro-2-(phenylsulfonyl)-8-isoquinolinyl]-1-piperazinyl]ethyl]-1,2,3,4-tetrahydro-, (3R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



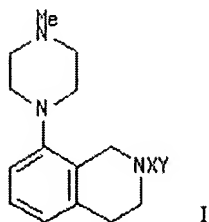
RE.CNT 20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 7 CAPLUS COPYRIGHT 2006 ACS on STN
Full Text

STN Columbus

AN 2004:1080868 CAPLUS
 DN 142:38280
 TI Preparation of piperazinyltetrahydroquinolines as serotonin 5-HT1B antagonists.
 IN Lowe, John Adams, III
 PA Pfizer Products Inc., USA
 SO PCT Int. Appl., 23 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004108682	A2	20041216	WO 2004-IB1855	20040601
	WO 2004108682	A3	20050428		
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	RW:	BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
	US 2004254193	A1	20041216	US 2004-852726	20040524
PRAI	US 2003-477559P	P	20030611		
OS	MARPAT 142:38280				
GI					



AB Title compds. [I; X = CH₂, CH₂CH₂, CO, COCH₂, OCH₂, OCH₂CH₂; Y = alkyl, (substituted) Ph, naphthyl, heteroaryl], were prepd. Thus, 8-bromoisoquinoline and PhCH₂BVr were refluxed together for 6 h in EtOH to give an oil which was stirred with NaBH₃CN in MeOH for 5 days to give 59% N-benzyl-8-bromotetrahydroisoquinoline. This was heated with N-methylpiperazine, Pd(OAc)₂, BINAP, and NaOCMe₃ in PhMe at 120° for 16 h to give 84% N-benzyl-8-(4-methylpiperazin-1-yl)tetrahydroisoquinoline. The latter was hydrogenolyzed followed by acylation with 4-chlorophenylacetic acid in the presence of EDAC and N-hydroxybenzotriazole in CH₂Cl₂ at room temp. to give N-(4-chlorophenylacetyl)-8-(4-methylpiperazin-1-yl)tetrahydroisoquinoline. I showed 5-HT1B receptor affinity with IC₅₀ <0.60 μM.

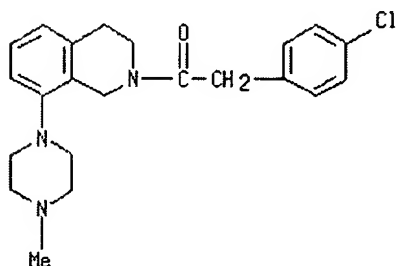
IT 807330-10-5P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(claimed compd.; prepn. of piperazinyltetrahydroquinolines as serotonin 5-HT1B antagonists)

STN Columbus

RN 807330-10-5 CAPLUS
 CN Isoquinoline, 2-[(4-chlorophenyl)acetyl]-1,2,3,4-tetrahydro-8-(4-methyl-1-piperazinyl)- (9CI) (CA INDEX NAME)



L4 ANSWER 3 OF 7 CAPLUS COPYRIGHT 2006 ACS on STN

Full Text

AN 2004:756620 CAPLUS

DN 141:277505

TI Preparation of 2,5- and 2,6-substituted tetrahydroisoquinolines as 5-HT6 modulators, in particular selective 5-HT6 antagonists, for treating CNS disorders

IN Putman, David George

PA F. Hoffmann-La Roche A.-G., Switz.

SO PCT Int. Appl., 59 pp.

CODEN: PIXXD2

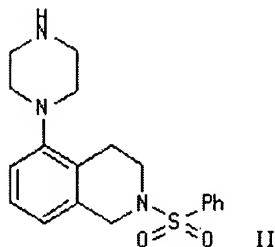
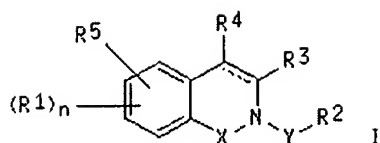
DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004078176	A1	20040916	WO 2004-EP1751	20040223
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	CA 2517146	AA	20040916	CA 2004-2517146	20040223
	EP 1601358	A1	20051207	EP 2004-713547	20040223
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	US 2004180874	A1	20040916	US 2004-791578	20040302
PRAI	US 2003-451516P	P	20030303		
	WO 2004-EP1751	W	20040223		
OS	MARPAT 141:277505				
GI					

APPs



AB Title compds. I [wherein n = 0-3; X = CH₂ and derivs.; C(:O); Y = SO₂ when X = CH₂ and derivs., and Y = (CH₂)_p and derivs. when X = C(:O); p = 1-3; each R₁ = independently halo, hetero/halo/alkyl, OH and derivs., NO₂, CN, S(O)_qH and derivs., NH₂ and derivs., CONH₂ and derivs., SO₂NH₂ and derivs., etc.; q = 0-2; R₂ = hetero/aryl, cycloalkyl; R₃, R₄ = independently H, alkyl; R₅ = 5- to 7-membered nitrogen-contg. heterocycle attached to the 5 or 6 position of the isoquinoline ring; their pharmaceutically acceptable salts or prodrugs] were prepd. as 5-HT₆ modulators for the treatment of central nervous system (CNS) disorders. Claimed are 5-HT₆ agonists useful for treating CNS diseases. A 5-step synthesis is given for tetrahydroisoquinoline salt II•TFA. In an in vitro radioligand binding study, I were found selective 5-HT₆ antagonists. Selected I showed high affinity for 5-HT₆ receptor with pK_i values in the range 9.28-10.37 in a radioligand assay. Thus, I are useful in treating mental disorders.

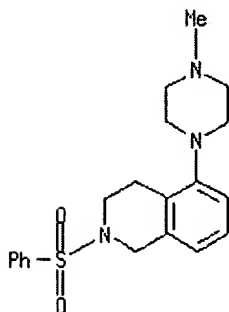
IT 757966-35-1P, 2-Benzenesulfonyl-5-(4-methylpiperazin-1-yl)-1,2,3,4-tetrahydroisoquinoline

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(5-HT₆ antagonist; prepn. of tetrahydroisoquinolines as 5-HT₆ modulators for treating CNS disorders)

RN 757966-35-1 CAPLUS

CN Isoquinoline, 1,2,3,4-tetrahydro-5-(4-methyl-1-piperazinyl)-2-(phenylsulfonyl)- (9CI) (CA INDEX NAME)



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RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 4 OF 7 CAPLUS COPYRIGHT 2006 ACS on STN

Full Text

AN 2003:356439 CAPLUS

DN 138:368779

TI Preparation of isoquinolines as 5-HT antagonists for treatment of
psychiatric disorders

IN Angst, Christof; Haeberlein, Markus; Hill, Daniel; Jacobs, Robert; Moore,
Gary; Pierson, Edward; Shenvi, Ashokkumar Bhikkappa

PA Astrazeneca AB, Swed.

SO PCT Int. Appl., 139 pp.

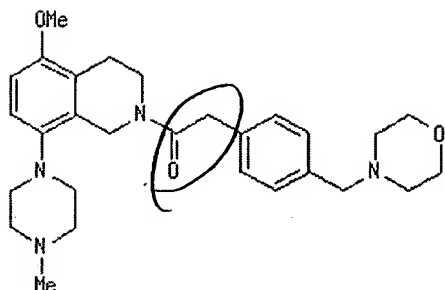
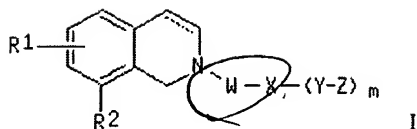
CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003037887	A1	20030508	WO 2002-SE1988	20021101
	WO 2003037887	C1	20050317		
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	GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,				
	LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,				
	PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,				
	UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW:				
	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,				
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	FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF,				
	CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	CA 2464342	AA	20030508	CA 2002-2464342	20021101
	EP 1451172	A1	20040901	EP 2002-780244	20021101
	R:				
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	BR 2002013778	A	20041109	BR 2002-13778	20021101
	JP 2005516896	T2	20050609	JP 2003-540168	20021101
	ZA 2004003240	A	20050407	ZA 2004-3240	20040429
	NO 2004002154	A	20040729	NO 2004-2154	20040525
PRAI	SE 2001-3644	A	20011101		
	WO 2002-SE1988	W	20021101		
OS	MARPAT 138:368779				
GI					



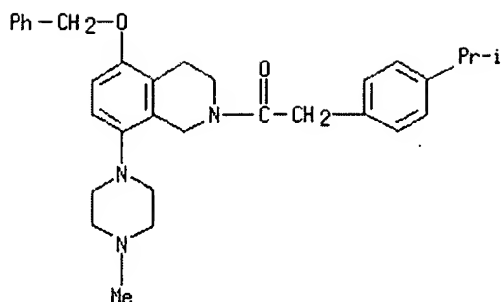
AB Title compds. I [wherein W = CO, CONRa, NRaCO, CO(CH₂)_nNRaCO, CSNRa, COCH₂O, SO₂NRa, NRaSO₂, CH₂NRa, COCH₂, CH₂CO, or 5-membered heterocyclyl; X = (un)substituted aryl or heterocyclyl; Y = bond, CH₂, O, S, SO, CO, SO₂, NRb, or NRbSO₂; Z = Rb, CO₂Ra, CON(Ra)₂, NHRb, alkyl-N(Ra)₂, SO₂Rc, or (un)substituted aryl(alkyl) or heterocyclyl; R1 = halo, alkyl, ORa, SOpRa, N(Ra)₂, or CN; R2 = aryl or heterocyclyl(carbonyl); Ra = H or (un)substituted alkyl; Rb = H, alkyl(sulfanyl), alkanoyl, aryl(alkyl), or arylalkoxyalkyl; Rc = alkyl, aryl, or heterocyclyl; m = 0 or 1; n = 0-4; p = 0-2;] were prepd. as 5-HT₁B and 5-HT₁D antagonists (no data). For example, O-methylation of 5-hydroxyisoquinoline using NaOBu-t and PhMe₃NC1 in DMF (85%), followed by bromination with bromine in AcOH gave 5-methoxy-8-bromoisquinoline (47%). Substitution with N-methylpiperazine using NaOBu-t, BINAP, and tris(dibenzylideneacetone)dipalladium in PhMe and subsequent redn. with NaCNBH₃ and BF₃•Et₂O in MeOH gave 5-methoxy-8-(4-methylpiperazin-1-yl)-1,2,3,4-tetrahydroisoquinoline. Coupling of 4-(bromomethyl)phenylacetic acid with morpholine in the presence of K₂CO₃ in MeCN provided 4-(morpholinomethyl)phenylacetic acid. Amidation of the tetrahydroisoquinoline with the phenylacetic acid in DMF afforded II. I are useful for the treatment of psychiatric disorders including but not limited to depression, generalized anxiety, eating disorders, dementia, panic disorder, and sleep disorders (no data). The compds. may also be useful in the treatment of gastrointestinal disorders, motor disorders, endocrine disorders, vasospasm, and sexual dysfunction (no data).

IT 521313-56-4P, 1-[5-Benzyloxy-8-(4-methylpiperazin-1-yl)-3,4-dihydro-1H-isoquinolin-2-yl]-2-(4-isopropylphenyl)ethanone
 RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
 (5-HT antagonist; prepn. of isoquinolines as 5-HT₁B and 5-HT₁D antagonists for treatment of psychiatric disorders)

RN 521313-56-4 CAPLUS

CN Isoquinoline, 1,2,3,4-tetrahydro-2-[[4-(1-methylethyl)phenyl]acetyl]-8-(4-methyl-1-piperazinyl)-5-(phenylmethoxy)- (9CI) (CA INDEX NAME)

STN Columbus



RE.CNT 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 5 OF 7 CAPLUS COPYRIGHT 2006 ACS on STN

Full Text

AN 2002:575065 CAPLUS

DN 137:140776

TI Preparation of piperidinyl and piperazinyl amino acid derivatives as melanocortin receptor agonists

IN Backer, Ryan Thomas; Briner, Karin; Doecke, Christopher William; Fisher, Matthew Joseph; Kuklish, Steven Lee; Mancuso, Vincent; Martinelli, Michael John; Mullaney, Jeffrey Thomas; Xie, Chaoyu

PA Eli Lilly and Company, USA

SO PCT Int. Appl., 263 pp.

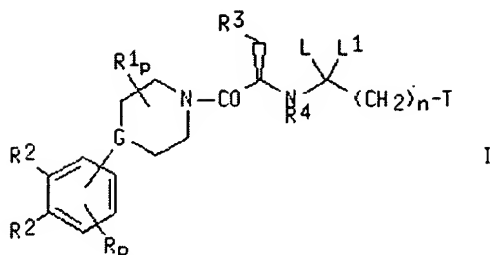
CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

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	EP 1368339	A1	20031210	EP 2002-701923	20020123
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	JP 2004521117	T2	20040715	JP 2002-559409	20020123
	US 2004058936	A1	20040325	US 2003-466249	20030711
PRAI	US 2001-263595P	P	20010123		
	WO 2002-US516	W	20020123		
OS	CASREACT 137:140776; MARPAT 137:140776				
GI					



I

AB The invention relates to melanocortin receptor (MC-R) agonists I [G = CR1 or N; LL1 = H2 or oxo; T = isoquinolinyl or tetrahydro deriv., isoindolinyl, or piperazinyl; n = 0-8; R = H, OH, CN, NO2, halo, alkyl, acyl, etc.; R1 = H, alkyl, alkylcarbamoyl, (D)phenyl, (D)cycloalkyl, or oxo (unless amide is formed); p = 0-4; CR2CR2 is a 5- or 6-membered carbocycle optionally substituted by 1-3 groups R; R3 = (un)substituted aryl or thienyl; R4 = H, alkyl, acyl, cycloalkyl, or alkoxyalkyl], or their pharmaceutically-acceptable salts or stereoisomers, which are useful in the treatment of obesity, diabetes, and male and/or female sexual dysfunction. Comps. I comprise three domains, i.e., a piperidino or piperazinyl fragment, an amino acid, and a radical CLL1(CH2)n-T. Thus, 1,2,3,4-tetrahydroisoquinoline-3-carboxylic acid [1-(4-chlorobenzyl)-2-[4-[2-(methylsulfonyl)-1,2,3,4-tetrahydroisoquinolin-8-yl]piperazin-1-yl]-2-oxoethyl]amide (claimed compd.) was prepd. via acylation of the piperazine moiety.

IT 444619-63-0P

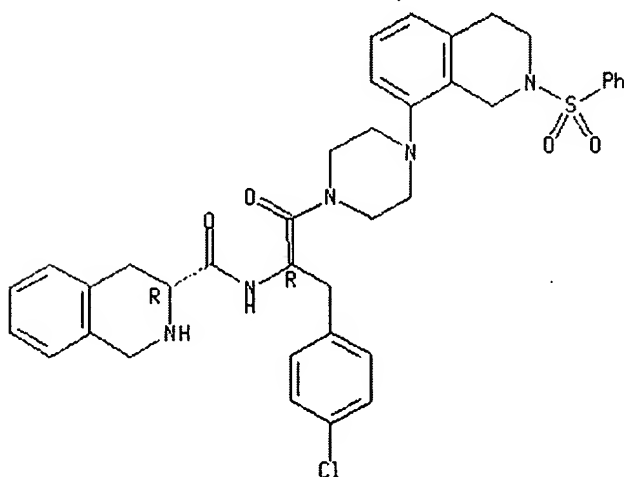
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of piperidiny and piperazinyl amino acid derivs. as melanocortin receptor agonists)

RN 444619-63-0 CAPLUS

CN 3-Isoquinolinecarboxamide, N-[(1R)-1-[(4-chlorophenyl)methyl]-2-oxo-2-[4-[1,2,3,4-tetrahydro-2-(phenylsulfonyl)-8-isoquinolinyl]-1-piperazinyl]ethyl]-1,2,3,4-tetrahydro-, (3R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RE.CNT 6

THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

STN Columbus

L4 ANSWER 6 OF 7 CAPLUS COPYRIGHT 2006 ACS on STN

Full Text

AN 2000:742072 CAPLUS

DN 133:309907

TI Preparation of nitrogen-containing heterocyclic compounds and benzamide compounds as hypolipidemics and antiarteriosclerotics

IN Ohkura, Naoto; Hiraiwa, Yukiko; Matsushima, Tetsuya; Sasaki, Kazue; Yamamoto, Takehiro; Shiotani, Masaharu; Suzuki, Shigeki; Nakatani, Yuuko; Kuroda, Chizuko; Nagasawa, Mieko; Katano, Kiyoaki

PA Meiji Seika Kaisha, Ltd., Japan

SO PCT Int. Appl., 284 pp.

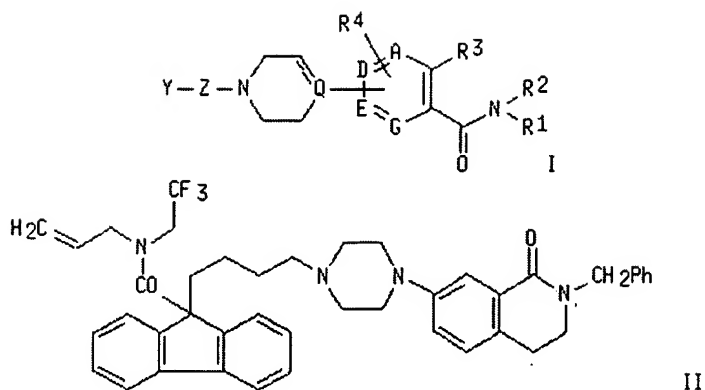
CODEN: PIXXD2

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000061556	A1	20001019	WO 2000-JP2329	20000410
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	RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	CA 2369103	AA	20001019	CA 2000-2369103	20000410
	BR 2000009650	A	20020102	BR 2000-9650	20000410
	EP 1180514	A1	20020220	EP 2000-915465	20000410
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	AU 779550	B2	20050127	AU 2000-36759	20000410
	US 6777414	B1	20040817	US 2001-958296	20011005
	US 2004224959	A1	20041111	US 2004-868006	20040616
PRAI	JP 1999-102559	A	19990409		
	JP 1999-118490	A	19990426		
	JP 1999-119043	A	19990427		
	WO 2000-JP2329	W	20000410		
	US 2001-958296	A3	20011005		
OS	MARPAT 133:309907				
GI					



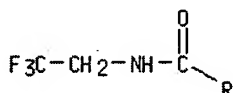
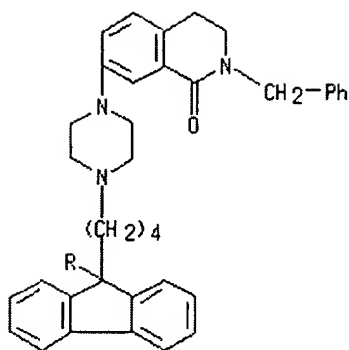
AB The title compds. [I; R1 and R2 represent each (un)substituted C1-6 alkyl or alkoxy, C3-8 cycloalkyl, Ph, C2-6 alkenyl or alkynyl, 5- or 6-membered ring (un)satd. heterocyclyl; R3 and R4 represent each hydrogen, (un)substituted C1-6 alkyl, halo, OH, cyano, C2-5 alkoxy-carbonyl, C1-6 alkoxy, or CO₂H; or R2 and R3 may be bonded to each other to form (CH₂)_m, N:CH, CH:N, or (C1-6 alkyl)-C:N; wherein m is 1 or 2; A, D, E and G represent each C, or one of A, D, E and G represents N and the remainders represent C; Q represents N or C; Y represents a group represented by general formula Q1 (wherein X represents hydrogen, CONR₅R₆, etc.; R₈ represents nil or a bond, O, etc.; and R₉ and R₁₀ represent each hydrogen, alkyl, etc.); and Z represents (CH₂)_n, O(CH₂)_i, or CONH(CH₂)_i; wherein n is 0-6; i is 1-6] are prepd. These compds. have an effect of inhibiting the biosynthesis of triglycerides in the liver and an effect of inhibiting the secretion of apolipoprotein B-contg. lipoproteins from the liver (the latter effect being particularly excellent), without showing the side effect of fat accumulation in the liver, and are useful in treating and preventing hyperlipemia, arteriosclerotic diseases, and pancreatitis. Thus, to a soln. of 2-benzyl-7-[4-[4-[9-(2,2,2-trifluoroethylcarbamoyl)-9H-fluorene-9-yl]butyl]piperazin-1-yl]-3,4-dihydro-2H-isoquinolin-1-one in PhMe were added NaOH, K₂CO₃, tetrabutylammonium hydrogen sulfate, and allyl bromide and the resulting mixt. was stirred at 60° overnight to give title compd. (II). II in vitro inhibited the secretion of apolipoprotein B by 89% and the biosynthesis of triglycerides by 89% in HepG2 cells. Tablet and capsule formulations were also described.

IT 301666-52-4P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of nitrogen-contg. heterocyclic compds. and benzamide compds. as hypolipidemics and antiarteriosclerotics and inhibitors of apolipoprotein B-contg. lipoproteins and biosynthesis of triglycerides)

RN 301666-52-4 CAPLUS

CN 9H-Fluorene-9-carboxamide, 9-[4-[4-[1,2,3,4-tetrahydro-1-oxo-2-(phenylmethyl)-7-isoquinolinyl]-1-piperazinyl]butyl]-N-(2,2,2-trifluoroethyl)- (9CI) (CA INDEX NAME)



RE.CNT 27 THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 7 OF 7 CAPLUS COPYRIGHT 2006 ACS on STN

Full Text

AN 1998:793126 CAPLUS

DN 130:52434

TI Prepn. of nitrogenous heterocyclic compounds as hyperlipemia remedies

IN Ohkura, Naoto; Tsuruoka, Takashi; Usui, Takayuki; Hiraiwa, Yukiko;
Matsushima, Tetsuya; Shiotani, Masaharu; Niizato, Tetsutaro; Nakatani,
Yuuko; Suzuki, Shigeki; Kuroda, Chidsuko; Katano, Kiyooki

PA Meiji Seika Kaisha, Ltd., Japan; et al.

SO PCT Int. Appl., 194 pp.

CODEN: PIXXD2

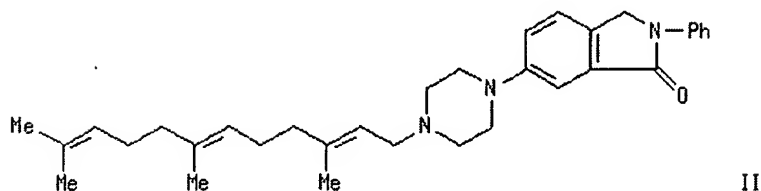
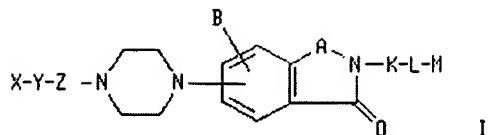
DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9854135	A1	19981203	WO 1998-JP2411	19980601
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	RW:	GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG			
	CA 2291630	AA	19981203	CA 1998-2291630	19980601
	AU 9875482	A1	19981230	AU 1998-75482	19980601
	EP 999208	A1	20000510	EP 1998-923066	19980601
	R:	DE, ES, FR, GB, IT			
	US 6417362	B1	20020709	US 1999-424708	19991130
	US 2002156276	A1	20021024	US 2002-127491	20020423
	US 6583144	B2	20030624		
PRAI	JP 1997-141410	A	19970530		
	WO 1998-JP2411	W	19980601		
	US 1999-424708	A3	19991130		
OS	MARPAT 130:52434				
GI					

STN Columbus



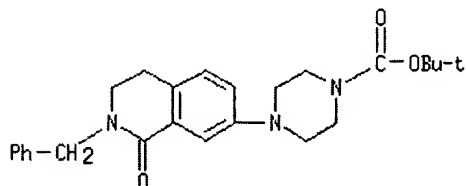
AB The title compds. [I; A = CR₁R₂(CH₂)_i; (wherein R₁ and R₂ each represents hydrogen or alkyl, i = 0-1), CH:CH, OCH₂, or S(O)_jCH₂ (wherein j = 0-2); B = hydrogen or halogen; X = CR₃R₄R₅, NR₆R₇, (CH₂CH:C(CH₃)CH₂)_pCH₂CH:C(CH₃)₂, alkyl, cycloalkyl, Ph, cinnamyl, or heteroaryl; Y = (CH₂)_q, CH:CH, NR₈, oxygen, or a bond; Z = carbonyl or a bond; K = alkylene or a bond; L = CH:CH or a bond; and M = hydrogen, alkyl, cycloalkyl, Ph, heterocycle, biphenyl, or diphenylmethyl; p = 0-2; q = 1-6; R₃-R₅ = hydrogen, phenyl; R₆-R₇ = hydrogen, Ph, benzyl; R₈ = hydrogen, C1-6 alkyl] are prepd. I inhibit the biosynthesis of triglycerides in the liver and also inhibit the secretion of lipoproteins contg. apolipoprotein B from the liver. I are hence useful for the prevention/treatment of hyperlipemia (esp. hyper-VLDL-emia) and diseases caused thereby, such as arteriosclerotic diseases, e.g., myocardial infarct, and pancreatitis. Thus, title compd. (II) was prepd. by multi-step reactions and showed 56% and 90% inhibitory activity for apolipoprotein B and triglycerides. A formulation contg. I was also presented.

IT 217492-19-8P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of nitrogenous heterocyclic compds. as hyperlipemia remedies)

RN 217492-19-8 CAPLUS

CN 1-Piperazinecarboxylic acid, 4-[1,2,3,4-tetrahydro-1-oxo-2-(phenylmethyl)-7-isoquinolinyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> file caold

COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION

FULL ESTIMATED COST

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DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE	TOTAL
ENTRY	SESSION

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L5 0 L3

=> file reg

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FULL ESTIMATED COST	0.44	209.02
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	0.00	-5.25

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* the IDE default display format and the ED field has been added, *
* effective March 20, 2005. A new display format, IDERL, is now *
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*

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<http://www.cas.org/ONLINE/UG/regprops.html>

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L1 STRUCTURE UPLOADED

L2 1 S L1

L3 198 S L1 SSS FULL

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L4 7 S L3

FILE 'CAOLD' ENTERED AT 11:46:07 ON 18 FEB 2006

L5 0 S L3

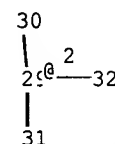
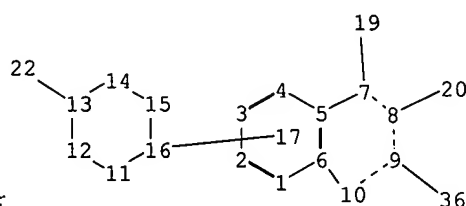
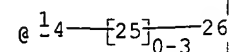
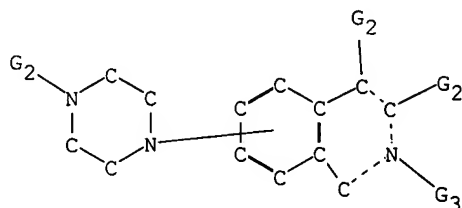
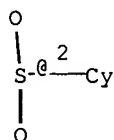
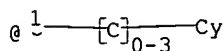
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ANSWER SET L3 HAS BEEN SAVED AS 'TEN791578/A'

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19 20 22 24 25 26 29 30 31 32 36

ring nodes :

1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16

chain bonds :

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ring bonds :

1-2 1-6 2-3 3-4 4-5 5-6 5-7 6-10 7-8 8-9 9-10 11-12 11-16
12-13 13-14 14-15 15-16

exact/norm bonds :

5-7 6-10 7-8 7-19 8-9 8-20 9-10 9-36 11-12 11-16 12-13 13-14
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exact bonds :

24-25

normalized bonds :

1-2 1-6 2-3 3-4 4-5 5-6

isolated ring systems :

containing 1 : 11 :

G2:C,H

G3:[*1],[*2]

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom
10:Atom 11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom
19:CLASS 20:CLASS 22:CLASS 24:CLASS 25:CLASS 26:Atom 29:CLASS
30:CLASS

31:CLASS 32:Atom 36:CLASS